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***** STN Columbus *****

FILE 'HOME' ENTERED AT 18:53:42 ON 20 JUN 2009

=> b ca

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FILE 'CA' ENTERED AT 18:54:00 ON 20 JUN 2009

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FILE COVERS 1907 - 18 Jun 2009 VOL 150 ISS 26

FILE LAST UPDATED: 18 Jun 2009 (20090618/ED)

REVISED CLASS FIELDS (/NCL) LAST RELOADED: Arp 2009

USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Feb 2009

CAS now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2008.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s apml or adiponectin

173 APM1
5472 ADIPONECTIN
L1 5556 APM1 OR ADIPONECTIN

=> s l1 and polyclonal antibody?

40047 POLYCLONAL
542111 ANTIBOD?
20073 POLYCLONAL ANTIBOD?
(POLYCLONAL(W)ANTIBOD?)
L2 15 L1 AND POLYCLONAL ANTIBOD?

=> s l2 and native(w)(apml or adiponectin)

156938 NATIVE
173 APM1

5472 ADIPONECTIN
 2 NATIVE(W) (APM1 OR ADIPONECTIN)
 L3 0 L2 AND NATIVE(W) (APM1 OR ADIPONECTIN)

=> s l2 and native
 156938 NATIVE
 L4 0 L2 AND NATIVE

=> s l2 and latex
 75640 LATEX
 L5 2 L2 AND LATEX

=> d ti ab 1-2

L5 ANSWER 1 OF 2 CA COPYRIGHT 2009 ACS on STN
 TI Determination of ***adiponectin*** in serum using a ***latex***
 particle-enhanced turbidimetric immunoassay with an automated analyzer
 AB ***Adiponectin*** is an adipose-derived hormone that plays a role in
 regulating metabolic processes such as fat partitioning and lipid and
 glucose metab. Quantification of ***adiponectin*** is useful for
 obtaining information on metabolic syndrome, but there is no rapid method
 to measure ***adiponectin*** for clin. use. The authors developed a
 rapid and sensitive ***latex*** particle-enhanced turbidimetric
 immunoassay (LTIA) using a ***latex*** bead-immobilized anti-
 adiponectin ***polyclonal*** ***antibody***. The assay
 was performed on a Hitachi H7170 analyzer and evaluated for validity as a
 method to quantitate ***adiponectin***, in parallel with the ELISA.
 Diln. tests using LTIA showed linearity from 0.25 to 30 .mu.g/mL.
 Within-run CV and total CV were obtained in the range 0.8-1.9% and
 1.1-2.0%, resp. No interference was obsd. in the testing of specimens
 contg. potentially interfering substances such as bilirubin,
 ditaurobilirubin, Hb triglyceride, rheumatoid factor, type IV collagen,
 fibronectin, and complement factor (Clq). A strong correlation between
 LTIA and ELISA was confirmed (n = 30, r = 0.990, y = 0.95x + 0.39). The
 LTIA assay is applicable to quantitating the serum concn. of
 adiponectin. This assay is more convenient and faster than ELISA
 and suitable for clin. routine anal.

L5 ANSWER 2 OF 2 CA COPYRIGHT 2009 ACS on STN
 TI ***Latex*** reagent for ***adiponectin*** analysis, and
 adiponectin analysis method
 AB A ***latex*** reagent for ***adiponectin*** anal. is provided,
 which comprises a suspension of ***latex*** particles carrying a
 substance (e.g., anti- ***adiponectin*** ***polyclonal***
 antibody) capable of specifically binding with
 adiponectin.
 . Also provided is a method for ***adiponectin*** anal., which
 comprises: (1) a step for obtaining a biol. liq. potentially contg.
 adiponectin; and (2) a step for contacting the biol. liq.
 obtained
 in the step (1) as it is with a suspension of ***latex*** particles
 carrying a substance capable of specifically binding with
 adiponectin, and optically analyzing the resultant mixt. to det.
 the degree of agglutination of the ***latex*** particles. According
 to this ***latex*** reagent for ***adiponectin*** anal. and this
 adiponectin anal. method, it is not required to dil. or pretreat

biol. liq. as a test sample beforehand, and the anal. is rapidly and conveniently performed without limiting a measurement facility.

=> d all 1-2

L5 ANSWER 1 OF 2 CA COPYRIGHT 2009 ACS on STN

AN 145:391809 CA <<LOGINID:20090620>>

ED Entered STN: 02 Nov 2006

TI Determination of ***adiponectin*** in serum using a ***latex*** particle-enhanced turbidimetric immunoassay with an automated analyzer

AU Nishimura, Ayako; Sawai, Tokio

CS Division of Research and Development, Mitsubishi Kagaku Iatron Inc.,

1460-6 Mitodai, Tako-machi, Katori-gun, Chiba-ken, 289-2247, Japan

SO Clinica Chimica Acta (2006), 371(1-2), 163-168

CODEN: CCATAR; ISSN: 0009-8981

PB Elsevier Ltd.

DT Journal

LA English

CC 9-10 (Biochemical Methods)

AB ***Adiponectin*** is an adipose-derived hormone that plays a role in regulating metabolic processes such as fat partitioning and lipid and glucose metab. Quantification of ***adiponectin*** is useful for obtaining information on metabolic syndrome, but there is no rapid method to measure ***adiponectin*** for clin. use. The authors developed a rapid and sensitive ***latex*** particle-enhanced turbidimetric immunoassay (LTIA) using a ***latex*** bead-immobilized anti-***adiponectin*** polyclonal ***antibody***. The assay was performed on a Hitachi H7170 analyzer and evaluated for validity as a method to quantitate ***adiponectin***, in parallel with the ELISA. Dilm. tests using LTIA showed linearity from 0.25 to 30 .mu.g/mL. Within-run CV and total CV were obtained in the range 0.8-1.9% and 1.1-2.0%, resp. No interference was obsd. in the testing of specimens contg. potentially interfering substances such as bilirubin, ditauobilirubin, Hb triglyceride, rheumatoid factor, type IV collagen, fibronectin, and complement factor (Clq). A strong correlation between LTIA and ELISA was confirmed (n = 30, r = 0.990, y = 0.95x + 0.39). The LTIA assay is applicable to quantitating the serum concn. of ***adiponectin***. This assay is more convenient and faster than ELISA and suitable for clin. routine anal.

ST ***adiponectin*** detn serum ***latex*** particle turbidimetric immunoassay automated analyzer; metabolic syndrome ***adiponectin*** serum immunoturbidimetry automatic analyzer

IT Cytokines

RL: ANT (Analyte); ANST (Analytical study)

(***adiponectin*** ; detn. of ***adiponectin*** in serum using ***latex*** particle-enhanced turbidimetric immunoassay with automated analyzer)

IT Blood analysis

Human

Immunoturbidimetry

Metabolic disorders

(detn. of ***adiponectin*** in serum using ***latex*** particle-enhanced turbidimetric immunoassay with automated analyzer)

RE.CNT 24 THERE ARE 24 CITED REFERENCES AVAILABLE FOR THIS RECORD

RE

(1) Anon; JAMA 2001, V285, P2486

- (2) Arita, Y; Biochem Biophys Res Commun 1999, V257, P79 CA
- (3) Arita, Y; Circulation 2002, V105, P2893 CA
- (4) Bland, J; Lancet 1986, V1, P307 MEDLINE
- (5) Daimon, M; Diabetes Care 2003, V26, P2015 CA
- (6) Haluzik, M; Physiol Res 2004, V53, P123 CA
- (7) Hansa, T; Clin Chem 1997, V43, P109
- (8) Hotta, K; Arterioscler Thromb Vasc Med 2000, V20, P1595 CA
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- (11) Lindsay, R; Lancet 2002, V360, P57 CA
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- (16) Okamoto, Y; Horm Metab Res 2000, V32, P47 CA
- (17) Ouchi, N; Circulation 1999, V100, P2473 CA
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- (22) Pischon, T; JAMA 2004, V291, P1730 CA
- (23) Ryo, M; Circ J 2004, V68, P975 CA
- (24) Tsukinoki, R; Lipids Health Dis 2005, V4, P27

L5 ANSWER 2 OF 2 CA COPYRIGHT 2009 ACS on STN

AN 141:310247 CA <<LOGINID::20090620>>

ED Entered STN: 28 Oct 2004

TI ***Latex*** reagent for ***adiponectin*** analysis, and
adiponectin analysis method

IN Tachikawa, Tetsuya; Akamatsu, Suguru; Sawai, Tokio; Nishimura, Fumiko
PA Mitsubishi Kagaku Iatron, Inc., Japan; Otsuka Pharmaceutical Co., Ltd.
SO PCT Int. Appl., 26 pp.
CODEN: PIXXD2

DT Patent

LA Japanese

IC ICM G01N033-53

ICS G01N033-543

CC 9-10 (Biochemical Methods)

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2004086040	A1	20041007	WO 2004-JP4083	20040324
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
	RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
	AU 2004223553	A1	20041007	AU 2004-223553	20040324
	AU 2004223553	B2	20081204		
	CA 2520438	A1	20041007	CA 2004-2520438	20040324
	EP 1607742	A1	20051221	EP 2004-723044	20040324

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK
 US 20070037207 A1 20070215 US 2005-550324 20050923
 PRAI JP 2003-80763 A 20030324
 WO 2004-JP4083 W 20040324

CLASS

PATENT NO. CLASS PATENT FAMILY CLASSIFICATION CODES

PATENT NO.	CLASS	PATENT FAMILY CLASSIFICATION CODES
WO 2004086040	ICM	G01N0033-53
	ICS	G01N0033-543
	IPCI	G01N0033-53 [ICM,7]; G01N0033-543 [ICS,7]
	IPCR	G01N0033-53 [I,C*]; G01N0033-53 [I,A]; G01N0033-543 [I,C*]; G01N0033-543 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
AU 2004223553	ECLA	G01N0033/543D; G01N0033/68V
	IPCI	G01N0033-53 [I,C*]; G01N0033-53 [I,A]; G01N0033-543 [I,C*]; G01N0033-543 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
	IPCR	G01N0033-53 [I,C*]; G01N0033-53 [I,A]; G01N0033-543 [I,C*]; G01N0033-543 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
CA 2520438	ECLA	G01N0033/543D; G01N0033/68V
	IPCI	G01N0033-53 [ICM,7]; G01N0033-543 [ICS,7]
	IPCR	G01N0033-53 [I,C*]; G01N0033-53 [I,A]; G01N0033-543 [I,C*]; G01N0033-543 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
EP 1607742	ECLA	G01N0033/543D; G01N0033/68V
	IPCI	G01N0033-53 [ICM,7]; G01N0033-543 [ICS,7]
	IPCR	G01N0033-53 [I,C*]; G01N0033-53 [I,A]; G01N0033-543 [I,C*]; G01N0033-543 [I,A]; G01N0033-68 [I,C*]; G01N0033-68 [I,A]
US 20070037207	ECLA	G01N0033/543D; G01N0033/68V
	IPCI	G01N0033-53 [I,A]; G01N0033-551 [I,A]
	NCL	435/007.100; 436/524.000

AB A ***latex*** reagent for ***adiponectin*** anal. is provided, which comprises a suspension of ***latex*** particles carrying a substance (e.g., anti- ***adiponectin*** ***polyclonal*** ***antibody***) capable of specifically binding with ***adiponectin***

. Also provided is a method for ***adiponectin*** anal., which comprises: (1) a step for obtaining a biol. liq. potentially contg. ***adiponectin*** ; and (2) a step for contacting the biol. liq. obtained in the step (1) as it is with a suspension of ***latex*** particles carrying a substance capable of specifically binding with ***adiponectin*** , and optically analyzing the resultant mixt. to det. the degree of agglutination of the ***latex*** particles. According to this ***latex*** reagent for ***adiponectin*** anal. and this ***adiponectin*** anal. method, it is not required to dil. or pretreat a biol. liq. as a test sample beforehand, and the anal. is rapidly and conveniently performed without limiting a measurement facility.

ST ***adiponectin*** analysis ***latex*** agglutination reagent antibody

IT Antibodies and Immunoglobulins
 RL: ARG (Analytical reagent use); ANST (Analytical study); USES (Uses)
 (***adiponectin*** anal. method using ***latex*** agglutination

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immunoassay reagent)
IT Cytokines
  RL: ANT (Analyte); ANST (Analytical study)
    ( ***adiponectin*** ; ***adiponectin*** anal. method using
      ***latex*** agglutination immunoassay reagent)
IT Agglutination test
  ( ***latex*** ; ***adiponectin*** anal. method using
    ***latex*** agglutination immunoassay reagent)
IT ***Latex***
  (particles; ***adiponectin*** anal. method using ***latex***
    agglutination immunoassay reagent)
RE.CNT 4      THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
RE
(1) A&t Corp; JP 20014624 A 2001
(2) Otsuka Pharmaceutical Co Ltd; EP 1033134 A 1999 CA
(3) Otsuka Pharmaceutical Co Ltd; US 6461821 B 1999 CA
(4) Otsuka Pharmaceutical Co Ltd; WO 9921577 A 1999 CA

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=> d his

(FILE 'HOME' ENTERED AT 18:53:42 ON 20 JUN 2009)

FILE 'CA' ENTERED AT 18:54:00 ON 20 JUN 2009

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L1      5556 S APM1 OR ADIPONECTIN
L2      15 S L1 AND POLYCLONAL ANTIBOD?
L3      0 S L2 AND NATIVE(W) (APM1 OR ADIPONECTIN)
L4      0 S L2 AND NATIVE
L5      2 S L2 AND LATEX

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=> logoff y

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	31.85	32.07
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-3.12	-3.12

STN INTERNATIONAL LOGOFF AT 18:56:58 ON 20 JUN 2009